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B02 RHONE-POULENC SANTE

RHON 13.07.83 *EP -133-098-A 13.07.83-FR-011706 (13.02.85) C07k-07

New S-della-methylene subsid. synergistin derivs. - useful as intermediates for antibacterials

C85-017097 D/S: AT BE CH DE FR GB IT LI LU NL SE.

Synergistin derivs, of formula (1), their addn. salts with acids and N-bases, metal salts and (where appropriate) their isomers or mixts, are new:

B(2-P1, 2-S) 2

= H or Me, N;

Y = H or Me, N;

R = (a) H or OH; (b) NR₁R₂ or (c) halo, trimethylsilyloxy, dialkylphosphoryloxy, -OSO₂R₃ or -OCOR₄;

R₁ and R₂ = H, phenyl or pyridyl (opt.substd. by di(1-4C)-alkylamino), 1-10C alkyl (opt.substd. by OH, SH, COOH, pyridyl, anilino, alkylamino or dialkylamino (with at least one alkyl substd. by OH, SH, COOH or anilino)), 3-4C alkenyl or alkynyl. alkynyl;

or R_1 and R_2 together complete a 5- or 6-membered heterocycle opt. contg. another O, S or N (opt.aikyl substd.) atom;

R₃ = alkyl, CF₁, CCl₃ or phenyl (opt.substd. by halo, alkyl or NO₂);
R₄ = as R₃ or also alkylcarbonylmethyl, 2-(alkylcarbonyl)-ethyl, alkoxycarbonylmethyl, 2-(alkoxycarbonyl)ethyl or alkoxy;

all alkyl contain 1-4C.

MORE SPECIFICALLY = H, OH or NR,'R,';

 R_1' and $R_2' = H$, phenyl (opt.substd. by dialkylamino).

alkyl (opt.substd. by OH, SH, COOH, pyridyl, anilino, alkylamino or dialkylamino, with alkyl substd. by OH) or 3-4C alkynyl.

USE (I) are intermediates in the synthesis of water-soluble antibacterial synergistin derivs. (which are claimed in

PREPARATION

 R_1 and R_2 = 1-4C alkyl or together complete a heterocycle; X_1 and X_2 = alkoxy or substd_amino as defined above for NR_1R_2.

Reaction is pref. at around 20°C, esp. using tert. butoxy bis(dimethylamino)methane (!!) as reactant.

The prod_can_be_reacted

The prod, can be reacted (1) with an alkali borohydride in the presence of a strong organic acid to give (1; R = H);
(2) with another amine to exchange the NR,R, gp.;
(3) hydrolysed to give R = OH which is then reacted

with halogenating agent, R'-halo. (R' = trimethylsilyloxy. dialkylphosphoryloxy, OSO_2R_3 or $OCOR_4$).

EXAMPLE

A soln, of 46g, pristinamycin I_A in 460 cc. 1,2-dichloro-ethane was treated with 230 cc. (11) and the mixt, stirred for 18 hr. at 20°C. It was then diluted with 11, dichloromethane, washed 3 times with 0.4% aq. NH4Cl, dried and

The residue was triturated with 600cc, water, filtered and Ine residue was triturated with bucc, water, liltered and the filtrate coned, to dryness to give 41g, crude 56-dimethylaminomethylene pristinamycin [A. A 23.5g, sample of this was chromatographed to give 12g, pure material of m.pt. about 195°C. (76pp1251HDDwgNo0/0).

(F) ISR: US4355112 5.Jnl.Ref.

EP-133098-A

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Office européen des brevets

DEMANDE DE BREVET EUROPEEN

(1) Numéro de dépôt: 84401479.5

(5) Int. Cl.4: C 07 K 7/00

(22) Date de dépôt: 12.07.84

(30) Priorité: 13.07.83 FR 8311706

Date de publication de la demande: 13.02.85 Bulletin 85/7

(M) Etats contractants désignés: AT BE CH DE FR GB IT LI LU NL SE

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Nouveaux dérivés de synorgistines et leur préparation.

(i) Nouveaux dérivés de synergistines de formule (i) dans lequelle Y = H ou N(CH.); et R représente

a) soit H ou OH

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b) soit un radical de formule NR₁R₂ dans laquelle R₁ et R₂ = H, phényle, pyridyle (éventuellement substitués par dialcoylamino (1 à 4 C) ou alcoyle (1 à 10 C) (éventuellement substitué par OH, SH, COOH, anilino, alcoylamino ou dialcoylamino dont au moins l'une des parties alcoyle est substituée par OH, SH, COOH ou anilino) ou alcènyle (3 ou 4 C), alcynyle (3 ou 4 C) ou bien R₁ et R₂ forment ensemble un hétérocycle contenant éventuellement un autre hétéroatome tel que O, S ou N (éventuellement substitué par alcoyle)

c) soit un atome d'halogène, un radical triméthylsilyloxy, dialcoylphosphoryloxy ou un radical -OSO-Rs ou -OCORs, Rs étant alcoyle, trifluorométhyle, trichlorométhyle, phényle eventuellement substitué et Rs étant défini comme Rs ou un radical ocylelcoyle, alcoxycarbonylalcoyle ou alcoyloxy, ainsi que leurs sels et leur préparation.

Ces produits sont utiles comme intermédiaires de synthèse.

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